I hereby certify that this is being deposited with the United States Postal Service under 37 CFR 1.8 on the date indicated above and is addressed to the Assistant Commissioner for Patents, Box: SEQUENCE Patriog ton, Date of Deposit is MAY_

PATENT/DX0804K

TRADEMARK OFFICE IN THE UNITED STATES PASEN

In re application of:

PARHAM, et al.

Serial No.:

09/265,540

Filed: 08-MAR-1999

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HUMAN RECEPTOR PROTEINS;

METHODS

RELATED REAGENTS AND

Assistant Commissioner for Patents

Box: SEQUENCE Washington, D.C. 20231

F. Hamud Examiner:

Art Unit:

1646

MECENED 1714 OB 5000

CORRECTION OF SEQUENCE

SUBMISSION

Palo Alto, California 94304-1104

JECH CENTER 1600/2900

-MAY-2000

RECEIVED

JUN 08 2000

TECH CENTER 1600/2900

COMPLIANCE WITH REQUIREMENTS FOR PATENT APPLICATIONS CONTAINING NUCLEOTIDE SEQUENCE AND/OR AMINO ACID SEQUENCE **DISCLOSURES**

Sir:

In reply to receipt of a "Notice to Comply" dated 27-APR-00 (Paper No. 6) for the above-identified application, Applicants submit: (1) a substitute paper and write-protected computer readable copy of the Sequence Listing as required under 37 CFR §1.825(d); (2) a statement under 37 CFR §1.825(d) and (3) a copy of the "Error Report." Since 27-MAY-00 falls on a Saturday, the period for a timely response is extended to the following business day which is Tuesday, 30-MAY-00 because Monday, 29-MAY-00 was a United States Patent & Trademark Office holiday.

Remarks

The enclosed sequence submission corrects the previous sequence submission by uniformly applying the PATENTIN 2.0 "new" format throughout the sequence listing. To the best of Applicants' knowledge and belief, the informational content of 25 the sequence itself remains unchanged; only the presentation format has been changed. For example, in SEQ ID NO: 1 the numeric identifiers <220> to <223> were newly added to the sequence listing to explain the use of the nucleotide symbol "n."

Applicants enclose the submission on a write protected floppy diskette containing the substitute "Sequence Listing." The diskette complies with the requirements of 37 CFR §1.824 and is IBM PC compatible with a PC-DOS/MS-DOS operating system. If the diskette is damaged, please contact Applicants' representative and a replacement diskette will be provided. A paper readable copy of the substitute Sequence Listing on the diskette is attached hereto.

The informational content of the substitute paper and substitute computer readable copies of the sequence listing, submitted in accordance with 37 CFR §1.825(d) are believed to be the same and are believed to include no new matter since the informational content of the enclosed sequences should be the same as the informational content of the sequences submitted in priority documents.

The enclosed items are a bona fide effort to bring the present application into full compliance with the rules for sequence submissions. Applicants have invested over five extra hours of significant additional labor and care in preparing the present submission. Should there be any question regarding the sufficiency of this submission to comply with the rules for sequence submissions, Applicants respectfully request notification of any specific deficiencies and an opportunity to remedy them, under 37 CFR §1.135(c).

No fees should be due, however, if any are required, Applicants authorize the Commissioner to make any appropriate charges or credit any overpayments to DNAX Research Institute Deposit Account No. 04-1239.

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2000 Dated: MAY

Respectfully, submitted

Gerald P. Keleher Reg. No. 43,707

30 Enclosures and attachments:

One substitute write-protected diskette (CRM) Substitute paper copy of contents of diskette Copy of the Sequence Error Report

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DNAX Research Institute 901 California Avenue Palo Alto, California 94304-1104 TEL: (650) 852-9196

40 FAX: (650) 496-1200

Apple tion No.	09/	26515	40.
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MOTICE TO COMPLY WI REQUIREMENTS FOR PATENT . PLICATIONS CONTAINING MUCLEOTIDE SEQUENCE AND/OR AMINO ACID SEQUENCE DISCLOSURES

The nucleotide and/or amino acid sequence disclosure contained in this application does not comply with the requirements for such a disclosure as set forth in 37 CFR 1.821 1.825 for the following reason(s):

~	
THE P	1. This application clearly fails to comply with the requirements of 37 CFR 1.8
	1.825. Applicant's attention is directed to these regulations, published at 11.29, May 15, 1990 and at 55 FR 18230, May 1, 1990.
	2. This application does not contain, as a separate part of the disclosure on
	paper copy, a "Sequence Listing" as required by 37 CFR 1.821(c).
മ	3. A copy of the "Sequence Listing" in computer readable form has not been
	submitted as required by 37 CFR 1.821(e).
	4. A copy of the "Sequence Listing" in computer readable form has been submitte
	However, the content of the computer readable form does not comply with the requirements of 37 CFR 1.822 and/or 1.823, as indicated on the attached copy of marked-up "Raw Sequence Listing."
سا	5. The computer readable form that has been filed with this application has been
	found to be damaged and/or unreadable as indicated on the attached CRF Diskette Problem Report. A substitute computer readable form must be submitted as requiby37 CFR 1.825(d).
	6. The paper copy of the "Sequence Listing" is not the same as the computer
	readable form of the "Sequence Listing" as required by 37 CFR 1.821(e).

Applicant must provide:

Other: -

An initial or substitute computer readable form (CRF) copy of the "Sequence Listing"

An initial or substitute paper copy of the "Sequence Listing", as well as an amendment directing its entry into the specification

A statement that the content of the paper and computer readable copies are the s and, where applicable, include no new matter, as required by 37 CFR 1.821(e) or 1.821(f) or 1.821(g) or 1.825(b) or 1.825(d)

For questions regarding compliance with these requirements, please conta

For Rules Interpretation, call (703) 308-1123

For CRF submission help, call (703) 308-4212

For PatentIn software help, call (703) 557-0400

Please return a copy of this notice with your response.